Amendment dated September 22, 2006 Reply to Office Action of March 22, 2006

AMENDMENTS TO THE CLAIMS

Please amend the claims as follows:

1. (Currently amended) A method for separating and purifying a-nucleic acid from a

biological sample, comprising a step of:

adsorbing and desorbing said a-nucleic acid to and from a membrane of an organic

macromolecule which has a membrane thickness of 10 µm to 500 µm, wherein the organic

macromolecule comprises surface-saponified acetylcellulose.

2. (Original) The method according to claim 1, wherein the organic macromolecule

is an organic macromolecule having a hydroxyl group on surface thereof.

3. (Canceled)

4. (Original) The method according to claim 1, wherein the organic macromolecule

is surface-saponified triacetylcellulose.

(Currently Amended) The method according to <u>claim 1</u> elaim 3, wherein the

surface-saponification rate of acetylcellulose is 5% or higher.

6. (Currently Amended) The method according to claim 1 elaim 3, wherein the

surface-saponification rate of acetylcellulose is 10% or higher.

2

Application No. 10/621,329 Docket No.: 0649-0965P

Amendment dated September 22, 2006 Reply to Office Action of March 22, 2006

7. (Original) The method according to claim 2, wherein acetylcellulose is a porous

film.

8. (Original) The method according to claim 2, wherein acetylcellulose is a non-

porous film.

9. (Previously presented) The method according to claim 1, wherein the nucleic acid

is in a sample solution when being adsorbed to and desorbed from the membrane of organic

macromolecule which has a membrane thickness of 10 μm to 500 μm.

10. (Original) The method according to claim 9, wherein the sample solution is a

solution prepared by adding a water-soluble organic solvent to a solution obtained by treating a

cell- or virus-containing test sample with a nucleic acid-solubilizing reagent.

11. (Original) The method according to claim 10, wherein the nucleic acid-

solubilizing reagent is a guanidine salt, a surfactant and a proteolytic enzyme.

12. (Original) The method according to claim 1, comprising steps of:

adsorbing the nucleic acid to a membrane of the organic macromolecule;

washing the membrane using a nucleic acid-washing buffer; and

desorbing the nucleic acid adsorbed to the membrane by using a liquid capable of

desorbing the nucleic acid adsorbed to the membrane.

3

Application No. 10/621,329 Docket No.: 0649-0965P Amendment dated September 22, 2006

Reply to Office Action of March 22, 2006

13. (Original) The method according to claim 12, wherein the nucleic acid-washing

buffer is a solution containing 20 to 100 % by weight of methanol, ethanol, isopropanol or n-

propanol.

14. (Original) The method according to claim 12, wherein the liquid capable of

desorbing the nucleic acid adsorbed to the membrane is a solution having a salt concentration of

0.5 M or lower.

15. (Original) The method according to claim 1, wherein adsorption and desorption of

the nucleic acid is carried out by using an unit for separation and purification of nucleic acid in

which a container having at least two openings contains a membrane of the organic

macromolecule which has a membrane thickness of 10 µm to 500 µm.

16. (Original) The method according to claim 1, wherein adsorption and desorption of

the nucleic acid is carried out by using an unit for separation and purification of nucleic acid

which comprises (a) a membrane of the organic macromolecule which has a membrane thickness

of 10 μ m to 500 μ m, (b) a container having at least two openings and containing the membrane,

and (c) a pressure difference-generating apparatus connected to one opening of the container.

17. (Currently Amended) A method comprising steps of:

(a) preparing a sample solution containing a-nucleic acid by using a test sample and

Application No. 10/621,329 Docket No.: 0649-0965P Amendment dated September 22, 2006

Reply to Office Action of March 22, 2006

inserting one opening of a unit for separation and purification of nucleic acid, which comprises a

membrane of an organic macromolecule which has a membrane thickness of 10 μm to 500 μm, a

container having at least two openings and containing the membrane, and a pressure difference-

generating apparatus connected to one opening of the container, into said sample solution

containing the nucleic acid;

(b) sucking the sample solution containing the nucleic acid by making an inside of the

container in a reduced pressure condition by using the pressure difference-generating apparatus

connected to the other opening of the unit for separation and purification of nucleic acid, and

contacting the sample solution to a membrane of the organic macromolecule which has a

membrane thickness of 10 μ m to 500 μ m;

(c) making the inside of the container in a pressurized condition by using the pressure

difference-generating apparatus connected to the other opening of the unit for separation and purification of nucleic acid, and discharging the sample solution containing the sucked nucleic

(d) inserting one opening of the unit for separation and purification of nucleic acid

into the nucleic acid-washing buffer:

acid to an outside of the container:

(e) sucking the nucleic acid-washing buffer by making the inside of the container in

the reduced pressure condition by using the pressure difference-generating apparatus connected

to the other opening of the unit for separation and purification of nucleic acid, and contacting the

nucleic acid-washing buffer to the membrane:

(f) making the inside of the container in the pressurized condition by using the

pressure difference-generating apparatus connected to the other opening of the unit for separation

5

Application No. 10/621,329 Docket No.: 0649-0965P Amendment dated September 22, 2006

Reply to Office Action of March 22, 2006

and purification of nucleic acid, and discharging the sucked nucleic acid-washing buffer to the

outside of the container;

(g) inserting one opening of the unit for separation and purification of nucleic acid

into the liquid capable of desorbing the nucleic acid adsorbed to the membrane;

(h) making the inside of the container in the reduced pressure condition by using the

pressure difference-generating apparatus connected to the other opening of the unit for separation

and purification of nucleic acid, and sucking the liquid capable of desorbing the nucleic acid

adsorbed to the membrane to contact the liquid to the membrane; and

(i) making the inside of the container in the pressurized condition by using the

pressure difference-generating apparatus connected to the other opening of the unit for separation

and purification of nucleic acid, and discharging the liquid capable of desorbing the nucleic acid

adsorbed to the membrane to the outside of the container.

18. (Currently Amended) A method comprising steps of:

(a) preparing a sample solution containing the nucleic acid using a test sample and

injecting said sample solution containing the nucleic acid into one opening of a unit for

separation and purification of nucleic acid, which comprises a membrane of an organic

macromolecule which has a membrane thickness of 10 μ m to 500 μ m, a container having at least

two openings and containing the membrane, and a pressure difference-generating apparatus

connected to one opening of the container;

(b) making the inside of the container in the pressurized condition by using the

pressure difference-generating apparatus connected to said one opening of the unit for separation

Application No. 10/621,329

Amendment dated September 22, 2006 Reply to Office Action of March 22, 2006

and purification of nucleic acid, and discharging the injected sample solution containing the

nucleic acid from the other opening to contact the sample solution to a membrane of the organic

macromolecule which has a membrane thickness of 10 µm to 500 µm;

(c) injecting the nucleic acid-washing buffer into said one opening of the unit for

separation and purification of nucleic acid;

(d) making the inside of the container in the pressurized condition by using the

pressure difference-generating apparatus connected to said one opening of the unit for separation

and purification of nucleic acid, and discharging the injected nucleic acid-washing buffer from

said other opening to contact the nucleic acid-washing buffer to the membrane;

(c) injecting the liquid capable of desorbing the nucleic acid adsorbed to the

membrane into said one opening of the unit for separation and purification of nucleic acid; and

(f) making the inside of the container in the pressurized condition by using the

pressure difference-generating apparatus connected to said one opening of the unit for separation

and purification of nucleic acid, and discharging the liquid capable of desorbing the injected

nucleic acid from said other opening, so as to desorb the nucleic acid adsorbed to the membrane

and discharge the nucleic acid to the outside of the container.

19. (New) The method of claim 17, wherein the organic macromolecule comprises

surface-saponified acetylcellulose.

20. (New) The method of claim 18, wherein the organic macromolecule comprises

surface-saponified acetylcellulose.

7

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Docket No.: 0649-0965P